

Gnptab Cas9-KO Strategy

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Project Overview



Project Name

Gnptab

Project type

Cas9-KO

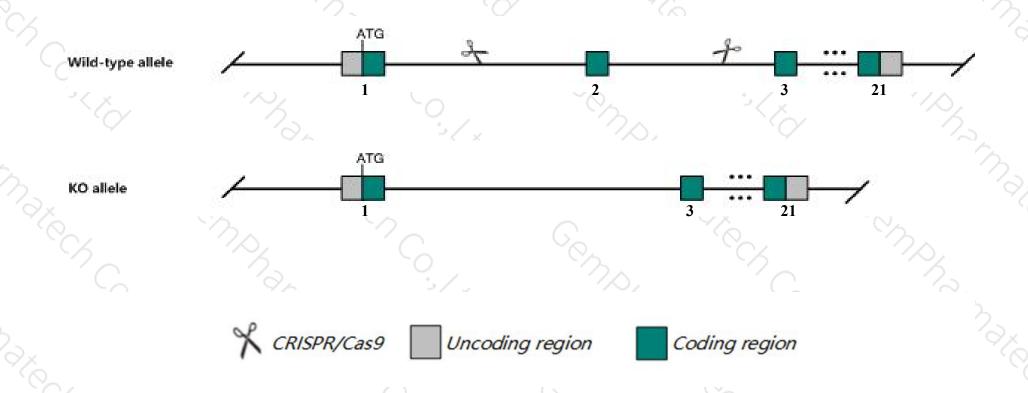
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Gnptab* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Gnptab* gene has 7 transcripts. According to the structure of *Gnptab* gene, exon2 of *Gnptab-201*(ENSMUST00000020251.9) transcript is recommended as the knockout region. The region contains 86bp coding sequence.

 Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Gnptab* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- > According to the existing MGI data, Homozygous mutations cause stunted growth, high lysosomal enzyme levels, skeletal defects, retinal degeneration and secretory cell lesions. Homozygotes for an ENU allele show skeletal and facial defects, altered enzymatic activities, lysosomal storage, Purkinje cell loss, ataxia and premature death.
- The *Gnptab* gene is located on the Chr10. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Gnptab N-acetylglucosamine-1-phosphate transferase, alpha and beta subunits [Mus musculus (house mouse)]

Gene ID: 432486, updated on 31-Jan-2019

Summary



Official Symbol Gnptab provided by MGI

Official Full Name N-acetylglucosamine-1-phosphate transferase, alpha and beta subunits provided by MGI

Primary source MGI:MGI:3643902

See related Ensembl:ENSMUSG00000035311

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as EG432486

Expression Ubiquitous expression in CNS E18 (RPKM 16.6), subcutaneous fat pad adult (RPKM 10.3) and 28 other tissuesSee more

Orthologs <u>human</u> all

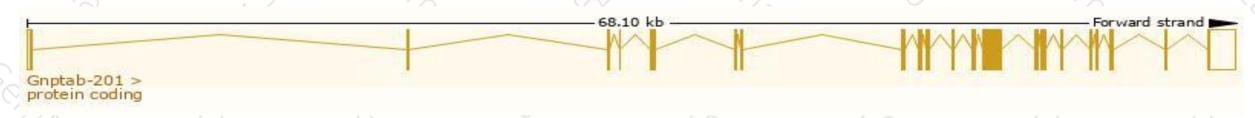
Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

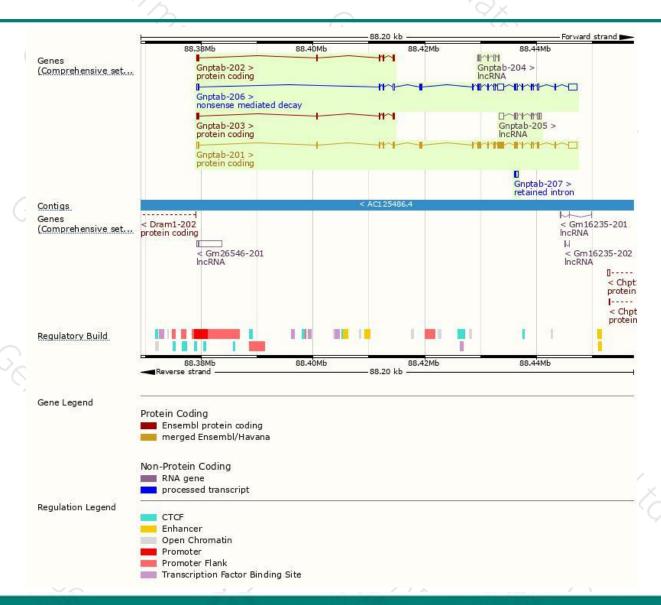
Transcript ID ENSMUST00000020251.9	bp 5390	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST00000020251.9	5390	100500				
		1235aa	Protein coding	CCDS24110	Q69ZN6	TSL:1 GENCODE basic APPRIS P1
ENSMUST00000127615.7	686	<u>174aa</u>	Protein coding		D3Z1C3	CDS 3' incomplete TSL:2
NSMUST00000130301.7	569	<u>166aa</u>	Protein coding	1/4/0	D3YXC6	CDS 3' incomplete TSL:5
NSMUST00000151273.7	5297	<u>56aa</u>	Nonsense mediated decay	767	D6RJ30	TSL:1
ENSMUST00000155306.1	536	No protein	Retained intron	-	-	TSL:3
ENSMUST00000141343.1	1856	No protein	IncRNA	(-)	-	TSL:1
NSMUST00000132738.1	726	No protein	IncRNA	1/20	2	TSL:3
	NSMUST00000130301.7 NSMUST00000151273.7 NSMUST00000155306.1	NSMUST00000130301.7 569 NSMUST00000151273.7 5297 NSMUST00000155306.1 536 NSMUST00000141343.1 1856	NSMUST00000130301.7 569 166aa NSMUST00000151273.7 5297 56aa NSMUST00000155306.1 536 No protein NSMUST00000141343.1 1856 No protein	NSMUST00000130301.7 569 166aa Protein coding NSMUST00000151273.7 5297 56aa Nonsense mediated decay NSMUST00000155306.1 536 No protein Retained intron NSMUST00000141343.1 1856 No protein IncRNA	NSMUST00000130301.7 569 166aa Protein coding - NSMUST00000151273.7 5297 56aa Nonsense mediated decay - NSMUST00000155306.1 536 No protein Retained intron - NSMUST00000141343.1 1856 No protein IncRNA -	NSMUST00000130301.7 569 166aa Protein coding - D3YXC6 NSMUST00000151273.7 5297 56aa Nonsense mediated decay - D6RJ30 NSMUST00000155306.1 536 No protein Retained intron - - NSMUST00000141343.1 1856 No protein IncRNA - -

The strategy is based on the design of *Gnptab-201* transcript, The transcription is shown below



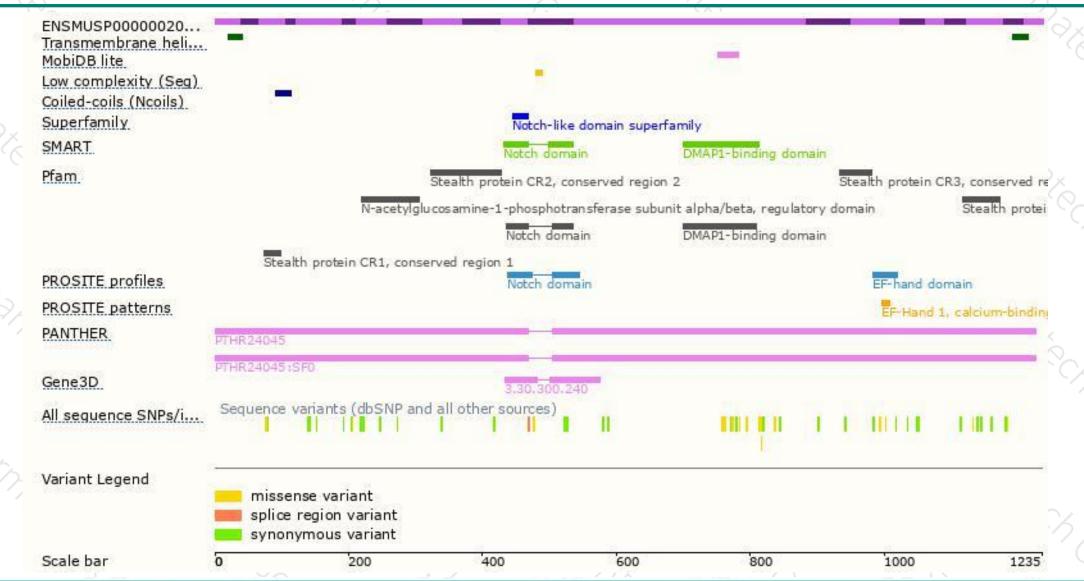
Genomic location distribution





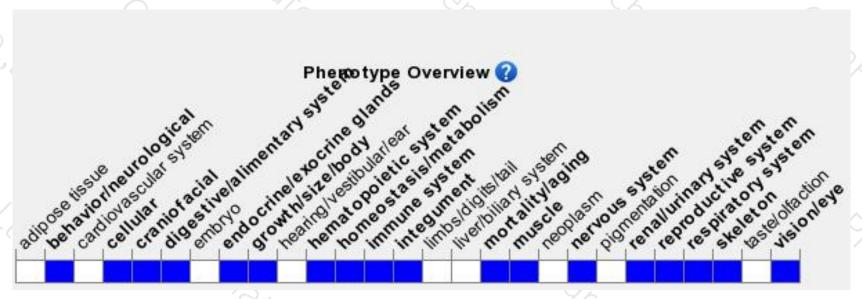
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

According to the existing MGI data, Homozygous mutations cause stunted growth, high lysosomal enzyme levels, skeletal defects, retinal degeneration and secretory cell lesions. Homozygotes for an ENU allele show skeletal and facial defects, altered enzymatic activities, lysosomal storage, Purkinje cell loss, ataxia and premature death.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





